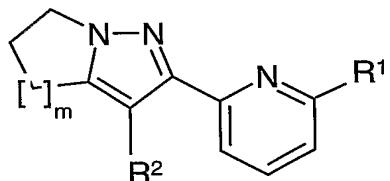


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5 WE CLAIM:

1. A compound of the formula:

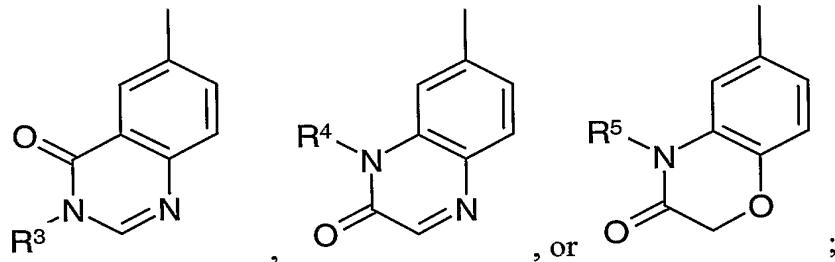


10

Formula I

wherein R¹ is hydrogen or (C₁-C₆) alkyl;

15 R² is selected from the group consisting of:



20 R³ is (C₁-C₆)alkyl, or (CH₂)_nX;

20

R⁴ is (C₁-C₆)alkyl, or (CH₂)_nX;

25 R⁵ is hydrogen, (C₁-C₆)alkyl, or (CH₂)_nX;

X is selected from the group consisting of a halogen, NR^aR^b,

N-morpholino, N-piperidine, N-pyrrolidine, or N-azepane;

n is an integer from 1-4;

m is an integer from 1-3;

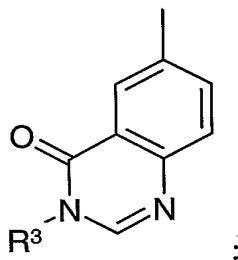
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5 R^a and R^b are each independently hydrogen or (C₁-C₆)alkyl;
and the pharmaceutically acceptable salts thereof.

2. A compound according to Claim 1 wherein:

R¹ is hydrogen or (C₁-C₆) alkyl;

10 R² is



R³ is (C₁-C₆)alkyl, or (CH₂)_nX;

X is selected from the group consisting of a halogen, NR^aR^b,

N-morpholino, N-piperidine, N-pyrrolidine, or N-azepane;

15 n is an integer from 1-4;

m is an integer from 1-3;

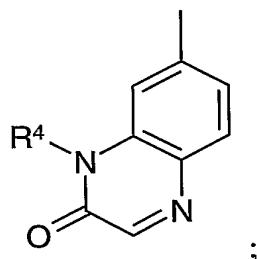
R^a and R^b are each independently hydrogen or (C₁-C₆)alkyl;

and the pharmaceutically acceptable salts thereof.

20 3. A compound according to Claim 1 wherein:

R¹ is hydrogen or (C₁-C₆) alkyl;

R² is



R⁴ is (C₁-C₆)alkyl, or (CH₂)_nX;

25 X is selected from the group consisting of a halogen, NR^aR^b,

N-morpholino, N-piperidine, N-pyrrolidine, or N-azepane;

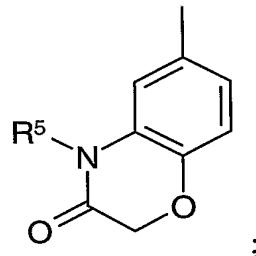
n is an integer from 1-4;

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5 m is an integer from 1-3;
R^a and R^b are each independently hydrogen or (C₁-C₆)alkyl;
and the pharmaceutically acceptable salts thereof.

4. A compound according to Claim 1 wherein:

10 R¹ is hydrogen or (C₁-C₆) alkyl;
R² is



R⁵ is hydrogen, (C₁-C₆)alkyl, or (CH₂)_nX;
X is selected from the group consisting of a halogen, NR^aR^b,
15 N-morpholino, N-piperidine, N-pyrrolidine, or N-azepane;
n is an integer from 1-4;
m is an integer from 1-3;
R^a and R^b are each independently hydrogen or (C₁-C₆)alkyl;
and the pharmaceutically acceptable salts thereof.

20 5. A compound according to Claims 1-4 wherein m is 1.

6. A compound according to Claims 1-5 wherein R³ is methyl.

25 7. A compound according to Claims 1-6 wherein R¹ is methyl.

8. A pharmaceutical formulation comprising a compound according to any one of Claims 1 to 7 in combination with a pharmaceutically acceptable diluent, excipient or carrier.

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5 9. A method of treating susceptible neoplasms in a patient in need thereof which comprises administering to said patient a therapeutically effective amount of a compound according to any one of Claims 1 to 7.

10 10. A method of treating fibrosis in a patient in need thereof which comprises administering to said patient a therapeutically effective amount of a compound according to any one of Claims 1-7.